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DESCRIPTION

SPHERICAL COMPOSITE COMPOSITION AND PROCESS FOR PRODUCING

SPHERICAL COMPOSITE COMPOSITION

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Technical Field

The present invention relates to a spherical composite composition and to a process of producing a spherical composite composition. Specifically, the invention relates to a spherical composite composition with high sphericity comprising a resin comprising unsaturated vinyl units and a magnetic material, which was difficult to produce effectively in the prior arts, and to a process of producing a spherical composite composition which is obtained by adding a magnetic material to an aqueous medium containing a resin dispersed therein, dispersing the material in the medium, and then forming the dispersion into particles by spray drying.

Background Art

20 In general, a resin polymerized in an aqueous medium is solidified as described in JP-B No. 1967-22684. Further, for a process of obtaining a spherical composite composition comprising a resin and a magnetic material, generally a resin and a magnetic material are melted and kneaded, crushed, classified, and formed into a spherical shape.

However, there are problems that the process requires two or more production steps and that the obtained composition has a broad distribution of the average particle diameters, and it is not always satisfactory in a viewpoint of productivity.

5 Especially, for a resin comprising acrylonitrile as a main component, there is a problem that thermal stability at the time of molding is so poor that the quality is lowered by thermal deterioration if melting and kneading are carried out with a magnetic material. In addition, for a resin

10 having high viscosity at the time of melting, a load occurs on the molding machine when melting and kneading are carried out with the magnetic material, which makes production impossible or worse.

Further, JP-A No. 1997-185184 describes a production process of dispersing a magnetic material in a resin.

15 However, since this process involves solution polymerization, the range of the molecular weight to be controlled is narrow, and thus the kind of the monomer is restricted by the kind of the solvent. In addition, there are other problems such

20 as a load in an environmental viewpoint by using the solvent, and the like.

[Patent Document 1] JP-B No. 1967-22684

[Patent Document 2] JP-A No. 1997-185184

Disclosure of the Invention

Problems to be Solved by the Invention

It is an object of the present invention to solve the above-mentioned problems and thus provide a spherical composite material comprising a resin comprising unsaturated vinyl units and a magnetic material, specifically a spherical composite composition produced by a simple and highly productive production process without the need of two or more production steps; and a process of producing the spherical composite composition. In particular, the invention provides a spherical composite composition obtained by adding a magnetic material to an aqueous medium containing a resin dispersed therein, dispersing the material in the medium, and then forming the dispersion into particles by spray drying; and a process of producing the spherical composite composition.

Means for Solving the Problems

The present inventors have studied extensively to solve such problems, and as a result, they have found that a spherical composite composition with high sphericity in the range of specific particle diameters is obtained by adding a magnetic material with a specific size to an aqueous medium containing a resin with specific the average particle

diameter dispersed therein, dispersing the material in the medium, and then forming the dispersion into particles by conducting spray drying under specific spray conditions to bind the resin with the magnetic material, which has led to
5 completion of the invention.

That is, the first aspect of the invention relates to a spherical composite composition which is made by adding (B) 5 to 1,000 parts by weight of a magnetic material having the longest length in two-dimensional projection of 0.01 to 50
10 μm , relative to 100 parts by weight of a resin comprising unsaturated vinyl units having (A-1) a glass transition temperature of 50 to 150°C and (A-2) a weight average molecular weight of 10,000 to 1,000,000, which is characterized in that the average particle diameter is 1 to
15 100 μm , and the sphericity is 0.7 to 1.

The second aspect of the invention relates to a process of producing a spherical composite composition which is the obtained by adding (B) 5 to 1,000 parts by weight of a magnetic material having the longest length in two-
20 dimensional projection of 0.01 to 50 μm , relative to 100 parts by weight of a resin dispersed in an aqueous medium comprising unsaturated vinyl units having (A-1) an average particle diameter of 0.01 to 1 μm , (A-2) a glass transition temperature of 50 to 150°C, and (A-3) a weight average

molecular weight of 10,000 to 1,000,000, dispersing the material in the medium, and then forming the dispersion into particles by spray drying, which is characterized in that the average particle diameter is 1 to 100 μm , and the 5 sphericity is 0.7 to 1.

The feature of the present production process is that after adding the magnetic material to an aqueous medium having a resin dispersed therein and dispersing the material in the medium, drying and particle formation are carried out 10 by spray drying at the same time. Therefore, the production process is very simple, has good productivity, and the obtained spherical composite composition has high sphericity.

Effects of the Invention

15 The spherical composite composition of the present invention has high sphericity, and can be suitably used in various applications such as a resin magnet, an electric wave absorption material, a magnetic shield material, and a developing agent and a toner carrier used in a developer of 20 the electric photograph process. Furthermore, the process of producing a spherical composite composition of the invention is a production process which makes it possible to produce a spherical composite composition with high sphericity comprising a resin and a magnetic material, which

was difficult to produce effectively in the prior arts,
whereby it is a production process which is very valuable
industrially.

5 Best Mode for Carrying Out the Invention

Hereinafter, the present invention will be explained in detail.

[Resin]

The first aspect of the invention is a spherical
10 composite composition with high sphericity comprising
specific unsaturated vinyl units and a magnetic material,
which can be effectively obtained by a production process of
adding a magnetic material to a resin dispersed in an
aqueous medium, dispersing the material in the medium, and
15 then forming the dispersion into particles by spray drying,
which is the second aspect of the invention.

In the invention, the resin may be subjected to
polymerization in an aqueous medium, or the powders after
the production may be dispersed in an aqueous medium.
20 However, considering the production of a resin having an
average particle diameter of 0.01 to 1 μm , polymerization in
an aqueous medium such as emulsion polymerization and
suspension polymerization is suitable in a viewpoint of
productivity, and further preferred is emulsion

polymerization. The smaller average particle diameter of the resin dispersed in the aqueous medium is suitable for dispersing the magnetic material. However, with the average particle diameter of less than 0.01 μm , it is very difficult

5 to produce a resin dispersed in an aqueous medium, and thus the practically preferable range is 0.01 μm or more.

Further, if the average particle diameter is too large, homogeneous dispersion is difficult, and thus it is preferably 1 μm or less. From such points, the average
10 particle diameter of the resin dispersed in an aqueous medium is 0.01 to 1 μm , and preferably 0.1 to 0.5 μm .

The above-mentioned resin comprising unsaturated vinyl units may be a homopolymer comprising one kind of the unsaturated vinyl units, or a copolymer comprising two or
15 more kinds of the unsaturated vinyl units. Further, as the unsaturated vinyl units, mention may be made of, for example, an unsaturated nitrile unit, a (meth)acrylic acid alkyl ester unit, an aromatic vinyl unit and the like.

As the monomer of the unsaturated nitrile unit, mention
20 may be made of, for example, acrylonitrile, methacrylonitrile, α -chloroacrylonitrile and the like, and furthermore, as the monomer of the (meth)acrylic acid alkyl ester unit, mention may be made of, for example, methyl (meth)acrylate, ethyl (meth)acrylate, propyl (meth)acrylate,

butyl (meth)acrylate and the like. As the monomer of the aromatic vinyl units, mention may be made of, for example, styrene, α -methyl styrene, vinyl toluene, vinyl xylene and the like. In addition, as the monomer of other 5 copolymerizable unsaturated vinyl units, mention may be made of, for example, vinyl ether, vinyl ester, α -olefin and the like. As the vinyl ester, mention may be made of, for example, vinyl acetate, vinyl propionate, vinyl butyrate and the like. As the vinyl ether, mention may be made of, for 10 example, methylvinyl ether, ethylvinyl ether, propylvinyl ether, butylvinyl ether, methylisopropenyl ether, ethylisopropenyl ether and the like. As the α -olefin, mention may be made of, for example, isobutene, 2-methyl-1-butene, 2-methyl-1-pentene, 2-methyl-1-hexene, 2-methyl-1-heptene, 2-methyl-1-octene, 2-ethyl-1-butene, 2-propyl-1-butene and the like.

Among the above-mentioned monomers of the unsaturated vinyl units, preferred is an acrylonitrile unit, a methacrylonitrile unit, a methyl (meth)acrylate unit, an 20 ethyl (meth)acrylate unit, a propyl (meth)acrylate unit, a butyl (meth)acrylate unit, a styrene unit, an α -methyl styrene unit or a vinyl toluene unit, and it is preferably contained in 30 to 100 % by weight, and preferably 50 to 100 % by weight in the resin. Furthermore, as used herein,

the (meth)acrylic acid alkyl ester means acrylic acid alkyl ester or methacrylic acid alkyl ester.

If the glass transition temperature of the above-mentioned resin is 50°C or higher, production without
5 generation of blocking at a spray part and aggregation upon recovery when the particles are formed becomes possible.

Considering the particle formation by spray drying, the glass transition temperature is from 50°C to 150°C, and preferably from 50°C to 110°C.

10 Further, the weight average molecular weight of the above-mentioned resin needs to be in the specific range, considering particle formation by spray drying. If the weight average molecular weight is 10,000 or more, the glass transition temperature does not become too low, and thus
15 production without generation of blocking at the spray part and aggregation upon recovery becomes possible. If the weight average molecular weight is 1,000,000 or less, the resin is melted by hot air at the time of particle formation, so it binds well to the magnetic material, which makes it
20 possible to obtain a composite composition with high sphericity. Considering such points, the weight average molecular weight is 10,000 to 1,000,000, and preferably 20,000 to 300,000.

[Magnetic material]

For the kind of the magnetic material used in the invention, mention may be made of, for example, a rare earth sintering type such as Nd-Fe-B series and Sm-Co series, ferrite sintering type such as Ba series, Sr series and La-
5 Co substitution series, soft ferrite type such as Mn-Zn series and Ni-Zn series, and miscellaneous, Al-Ni-Co series, Fe-Mn series, Fe-Cr-Co series, Sm-Fe-N series and the like. Further, the shape of the magnetic material is not especially limited, and the shape such as a circular
10 shape, a ring shape, an angular shape or a segment shape can be applied.

The smaller size of the magnetic material is more suitable for dispersion, but if it is too small, it is difficult to handle it. Thus, considering the average
15 particle diameter and the sphericity of the final composition, the longest length in two-dimensional projection is preferably 0.01 to 50 μm , and further preferably 0.1 to 10 μm . Further, the amount of the magnetic material to be added is 5 to 1,000 parts by weight,
20 preferably 10 to 800 parts by weight, relative to 100 parts by weight of the resin.

[Production of spherical composite resin]

The process of adding the magnetic material to the aqueous medium having a resin dispersed therein, and then

dispersing the material in the medium is not especially limited, and a general process can be used, such as a process wherein the magnetic material is added and then dispersed by a stirring blade, a process wherein dispersion 5 is carried out with a homogenizer, and other processes.

Total concentration of the resin and the magnetic material in the aqueous medium is preferably 10 to 85 % by weight, and further preferably 20 to 80 % by weight. If the total concentration is more than 10 % by weight, the concentration 10 is not low, and thus the productivity is not lowered. If the total concentration is less than 85 % by weight, the viscosity of the dispersion is not too high, and thus it is possible to prevent poor transport into the spray device or blocking at the spray part.

15 Furthermore, in the production of the spherical composite composition of the invention, a dispersing agent can be used in a combination with the aqueous medium. It is preferred that an anion surfactant be contained as the dispersing agent. Among them, at least one anion surfactant 20 selected from an alkylsulfuric acid ester salt, an alkylbenzene sulfonic acid salt, an alkylnaphthalene sulfonic acid salt, an alkylsulfosuccinic acid salt and a fatty acid salt is preferable as the dispersing agent.

Further, the amount of the dispersing agent to be used is

0.1 to 10 parts by weight, and preferably, 0.1 to 5 parts by weight, relative to 100 parts by weight of the monomer.

In the process of producing a spherical composite composition of the invention, the magnetic material is added 5 to an aqueous medium having a resin dispersed therein and dispersed in the medium in the above-mentioned manner, and then particle formation by spray drying is carried out. The spraying method in the spray drying is not especially limited, but it may be of a nozzle type, a disk type or 10 other types. Specifically, in the case of the nozzle type, a pressure nozzle type, a pressure two-fluid nozzle type, a two-fluid nozzle type or a four-fluid nozzle type can be employed, and in the case of the disk type, a pin form disk type, a vane form disk type, a Kestner form disk type and 15 the like. Among these, preferred is a pressure nozzle type, a two-fluid nozzle type, a pin form disk type or a vane form disk type, considering the long-run in production and the particle diameter distribution. The collection method is not especially limited, and a one-point collection method, a 20 two-point collection method or the like can be used.

Further, the heating source is not also especially limited, and an electricity type, a gas type, a vapor type or the like can be applied. For the method for contact with hot air, a co-current type, a counter-current type or a co-

current/counter-current type can be applied. Considering the kind and the concentration of the resin or the dispersion to be used, and the water content of the spherical composite composition after the spray drying, the 5 spray pressure in the case of the nozzle type and the disk rotation number in the case of the disk type are respectively controlled such that the average particle diameter of the obtained spherical composite composition is 1 to 100 μm and the sphericity is 0.7 to 1, and preferably 10 such that the average particle diameter is 5 to 70 μm and the sphericity is 0.75 to 1. If the disk rotation number in the case of the disk type is 3000 rpm or more, the sprayed drop is not so big that the average particle diameter of the spherical composite composition does not becomes too large 15 and at the same time, the spherical composite composition can be dried sufficiently. In addition, if the disk rotation number is 50,000 rpm or less, the sprayed drop is not so small that the average particle diameter of the spherical composite composition does not becomes too small. 20 Considering such points, the disk rotation number is preferably 3,000 to 50,000 rpm, and further preferably 5,000 to 20,000 rpm. In the particle formation by spray drying, it is difficult to produce a spherical composite composition having the average particle diameter of less than 1 μm ,

while if the average particle diameter is 1 μm or more, the spherical composite particle is not so small that it is easy to handle. Further, if the average particle diameter is 100 μm or less, the spherical composite particles can be 5 obtained in high yield after spray drying though it also depends on the size of the magnetic material to be blended, and the high sphericity of the obtained composite composition can be also maintained.

Furthermore, considering the productivity when 10 carrying out spray drying and particle formation at the same time, a specific range of drying condition is required. If the temperature of the inlet for hot air in the spray drying device is 100°C or higher, water may be evaporated during drying, and at the same time the resin may be melted 15 to form particles. If the temperature is the temperature which is the glass transition temperature of the resin plus 150°C or lower, continuous operation without generation of blocking by aggregation and solidification of the resin at the spray part is allowable. Furthermore, if the 20 temperature of the outlet for hot air in the spray drying device is 40°C or higher, the process of drying and particle formation is sufficiently carried out. If it is the temperature which is the glass transition temperature

of the resin plus 50°C or lower, it is possible to prevent the resin from adhering to the drying chamber and the like, which allows continuous operation. Considering such points,

5 when carrying out spray drying and particle formation at the same time, the temperature of the inlet for hot air in the spray drying device is from 100°C to the temperature which is the glass transition temperature of the resin plus 150°C, and temperature of the outlet for hot air in the spray drying device is from 40°C to the temperature which is the 10 glass transition temperature of the resin plus 50°C.

Preferably, the temperature of the inlet for hot air in the spray drying device spray is from 100°C to the temperature which is the glass transition temperature of the resin plus 100°C, and the temperature of the outlet for hot air in the 15 spray drying device is from 50°C to the temperature which is the glass transition temperature of the resin plus 20°C.

The composition obtained by the above-mentioned process is a spherical composite composition with high sphericity.

[Resin magnet]

20 The spherical composite composition with high sphericity of the invention as described above, can be used as a resin magnet. It can be used as a powder as it is, or can be used in various ways such as a powder paint, an applying agent as dispersed in water or a solvent, a molded

product after a molding process and the like. Further, the resin magnet can be used in a motor, an electric generator, a rotation controller, a magnet control, a speaker, an electromagnetic buzzer, a magnetic therapeutic machine, a
5 sensor, a magnet chuck and the like.

[Electric wave absorption material]

The spherical composite composition with high sphericity of the invention as described above, can be used as an electromagnetic wave absorption material. It can be
10 used as a powder as it is, or can be used in various ways such as a powder paint, an applying agent as dispersed in water or a solvent, a molded product after molding process and the like. In addition, the electric wave absorption material can be used for an inner wall material of an
15 electric wave dark room, a material for preventing reception interference by reflection of the broadcast wave, a material for preventing radar ghost by electric wave reflection and the like.

[Magnetic shield material]

20 The spherical composite composition with high sphericity of the invention as described above can be used as a magnetic shield material. It can be used as a powder as it is, or can be used in various ways such as a powder paint, an applying agent as dispersed in water or a solvent,

a molded product after molding process and the like. In addition, the magnetic shield material can be used in an internal magnetic shield material of an electrical/electronic device, a protecting shield material 5 of a monitor or a magnetic card, as a shield room and the like.

[Magnetic toner material used in the developer of electric photograph process]

The spherical composite composition with high 10 sphericity of the invention as described above can be used as a magnetic toner material for a one-component or two-component electric photograph. When it is used as a magnetic toner, a charge regulating agent, a surface treating agent, carbon black, a coloring agent, a wax and 15 the like may be further added in the inner part or on the surface.

[Toner carrier material used in the developer of electric photograph process]

The spherical composite composition with high 20 sphericity of the invention as described above can be used as a developing agent for a two-component electric photograph as mixed with the toner. When it is used as a toner carrier, it can be used as untreated, or can be used after carrying out surface treatment, heat treatment and the

like. A toner is that in which a coloring agent is dispersed in the binding resin. The amount of the toner to be contained is not especially limited. When the toner is used and consumed for development, it may be suitably added.

5 Usually, the content of the toner is about 10 to 100,000 parts by weight, relative to 100 parts by weight of the spherical composite product for the carrier.

Examples

10 Hereinafter, the present invention will be further illustrated with reference to Examples and Comparative Examples. The invention is not limited in any way to these examples as long as it is not beyond its gist.

Furthermore, "part" and "%" in Examples and Comparative Examples are intended to be based on weight. In addition, analysis and measurement described in Examples and Comparative Examples were carried out according to the following methods.

20 (1) Composition of the resin (% by weight)

The composition of carbon, hydrogen and nitrogen to be contained was measured by elemental analysis [manufactured by Yanagimoto Manufacturing, Co., Ltd., CHN CORDER, model:

MT-2], and this procedure was repeated three times and averaged to determine the composition of the resin.

5 (2) Average particle diameter of the resin in the aqueous medium (μm)

It was measured by HPPS which is a device for particle diameter measurement manufactured by Malvern Instruments Ltd. Furthermore, the average particle diameter is a value based on volume by dynamic light scattering.

10

(3) Glass transition temperature of the resin ($^{\circ}\text{C}$)

The temperature of the sample was elevated to 150°C under a nitrogen atmosphere, the sample was placed for 3 minutes at that temperature and then cooled to room temperature at a temperature lowering rate of $10^{\circ}\text{C}/\text{min}$.
15 Then, when measured using differential scanning calorimetry [manufactured by PERKIN-ELMER, model: DSC-7] at a temperature elevating rate of $5^{\circ}\text{C}/\text{min}$, the temperature of the cross point between the extension line of the baseline under the glass transition temperature, and the tangent line showing maximum slope from the first starting part of the peak to the top point of the peak, was taken as the glass
20 transition temperature.

(4) Weight average molecular weight of the resin

It was measured by gel permeation chromatography (hereinafter, referred to as GPC) using a mono-dispersed polystyrene standard sample as a standard.

5 [Measurement device and condition]

GPC: model: 150-C manufactured by Waters corporation

Column: model: 2 pieces of Shodex AD-80M/S
manufactured by Showa Denko Co.

Solvent: N,N-dimethylformamide (containing 0.1 % by

10 weight of lithium bromide)

Flow rate: 0.8 ml/min

Column temperature: 60°C

Sample concentration: 0.1 % by weight

Injection amount: 200 µl

15 Detector: Refractive index detection type.

(5) Longest length of the magnetic material (µm)

Using the photographs taken by an electron microscope, the longest lengths of 20 magnetic materials were measured
20 and then averaged.

(6) Productivity in particle formation

Particle formation was carried out for 2 hours continuously by a spray drying device. When nearly no

adhesion was found in the spray drying device and no solidified substance was generated in the disk of the spray part after completing the particle formation, the productivity was evaluated as AA; when much adhesion was 5 found, or a solidified substance was generated in the disk of the spray part after completing particle formation, the productivity was evaluated as BB; and when much amount of a solidified substance was generated in the disk of the spray part during particle formation so that operation was not 10 possible for more than 30 minutes, the productivity was evaluated as CC.

(7) Average particle diameter of the spherical composite composition (μm)

15 Using MT3000EX manufactured by MICRO TRACK CO., LTD., it was measured by a dry method. Furthermore, the average particle diameter is a value based on volume.

(8) Measurement of sphericity

20 Using the photographs taken by an electron microscope, (maximum diameters - minimum diameters) of 20 spherical composite compositions were measured, and then averaged. Then, using the value of the average particle diameter of

(6), the sphericity was calculated by the following calculation equation.

Value = [average particle diameter - (difference between the longest diameter and the shortest diameter in two-dimensional projection)]/average particle diameter.

[Production of the resin]

[Preparation Example 1]

15 parts of acrylonitrile, 5 parts of methyl acrylate, 10 0.407 part of sodium dioctylsulfosuccinate, 150 parts of water were put as raw materials of an initially added fraction to a stainless polymerization reactor, and the temperature was elevated to 60°C with stirring under a nitrogen atmosphere. The mixture was stirred for 30 minutes 15 as it was, an aqueous solution containing 0.08 part of potassium persulfate as a polymerization initiator was added to initiate polymerization.

Next, phosphoric acid was added to regulate pH as 3±0.3, and , polymerization was continued at 60°C while 20 continuously adding 60 parts of acrylonitrile, 20 parts of methyl acrylate, 1.6 parts of pentaerythritol tetrakis(β-mercaptopropionate), 1.627 parts of sodium dioctylsulfosuccinate and 85 parts of water as raw materials of subsequently added fraction.

During the addition, phosphoric acid was also continuously added for 5 hours starting from the initiation of polymerization, so polymerization was carried out for 6 hours with keeping the pH of the polymerization system as 5 3 ± 0.3 . When 8 hours passed from the initiation of polymerization, cooling was carried out to give a resin [A-1]. For this resin, the resin composition, the average particle diameter of the resin in the aqueous medium, the glass transition temperature and the weight average 10 molecular weight were measured according to the above-described methods. The results were as follows:

<Results of the measurement>

Resin composition: 75% acrylonitrile, 25% methyl acrylate

Average particle diameter of the resin in the aqueous medium: 0.22 μm

Glass transition temperature: 85°C

Weight average molecular weight: 120,000

20 [Preparation Example 2]

Polymerization was carried out in the same manner as in Preparation Example 1 except that 0.8 part of pentaerythritol tetrakis(β -mercaptopropionate) was used in

Preparation Example 1, to give a resin [A-2] having the
following results of the measurement:

<Results of the measurement>

Resin composition: 75% acrylonitrile, 25% methyl
5 acrylate

Average particle diameter of the resin in the aqueous
medium: 0.23 μm

Glass transition temperature: 86°C

Weight average molecular weight: 220,000

10

[Preparation Example 3]

Polymerization was carried out in the same manner as in
Preparation Example 1 except that 4 parts pentaerythritol
tetrakis(β -mercaptopropionate) were used in Preparation
15 Example 1, to give a resin [A-3] having the following
results of the measurement:

<Results of the measurement>

Resin composition: 75% acrylonitrile, 25% methyl
acrylate

20 Average particle diameter of the resin in the aqueous
medium: 0.20 μm

Glass transition temperature: 81°C

Weight average molecular weight: 30,000

[Preparation Example 4]

Polymerization was carried out in the same manner as in Preparation Example 1 while changing the composition of the initially added fraction and the subsequently added fraction 5 of the raw materials as below, to give a resin [A-4] having the following results of the measurement:

<Initially added fraction>

10 parts of acrylonitrile, 10 parts of methyl methacrylate

10 <Subsequently added fraction>

40 parts of acrylonitrile, 40 parts of methyl methacrylate

<Results of the measurement>

Resin composition: 50% acrylonitrile, 50% methyl
15 methacrylate

Average particle diameter of the resin in the aqueous medium: 0.22 μm

Glass transition temperature: 93°C

Weight average molecular weight: 130,000

20

[Preparation Example 5]

Polymerization was carried out in the same manner as in Preparation Example 1 while changing the composition of the initially added fraction and the subsequently added fraction

of the raw materials as below, to give a resin [A-5] having the following results of the measurement:

<Initially added fraction>

10 parts of acrylonitrile, 10 parts of styrene

5 <Subsequently added fraction>

40 parts of acrylonitrile, 40 parts of styrene

<Results of the measurement>

Resin composition: 50% acrylonitrile, 50% styrene

Average particle diameter of the resin in the aqueous

10 medium: 0.24 μm

Glass transition temperature: 95°C

Weight average molecular weight: 140,000

[Preparation Example 6]

15 Polymerization was carried out in the same manner as in Preparation Example 1 while changing the composition of the initially added fraction and the subsequently added fraction of the raw materials as below, to give a resin [A-6] having the following results of the measurement:

20 <Initially added fraction>

13 parts of acrylonitrile, 7 parts of butyl acrylate

<Subsequently added fraction>

52 parts of acrylonitrile, 28 parts of butyl acrylate

<Results of the measurement>

Resin composition: 65% acrylonitrile, 35% butyl acrylate

Average particle diameter of the resin in the aqueous medium: 0.22 μm

5 Glass transition temperature: 55°C

Weight average molecular weight: 130,000

[Preparation Example 7]

Polymerization was carried out in the same manner as in
10 Preparation Example 1 while changing the composition of the initially added fraction and the subsequently added fraction of the raw materials as below, to give a resin [A-7] having the following results of the measurement:

<Initially added fraction>

15 20 parts of acrylonitrile

<Subsequently added fraction>

80 parts of acrylonitrile

<Results of the measurement>

Resin composition: 100% acrylonitrile

20 Average particle diameter of the resin in the aqueous medium: 0.20 μm

Glass transition temperature: 100°C

Weight average molecular weight: 130,000

[Preparation Example 8]

Polymerization was carried out in the same manner as in Preparation Example 1 except that 0.2 part pentaerythritol tetrakis(β -mercaptopropionate) was used in Preparation Example 1, to give a resin [A-8] having the following results of the measurement:

<Results of the measurement>

Resin composition: 75% acrylonitrile, 25% methyl acrylate

Average particle diameter of the resin in the aqueous medium: 0.27 μm

Glass transition temperature: 90°C

Weight average molecular weight: 1,200,000

15 [Preparation Example 9]

Polymerization was carried out in the same manner as in Preparation Example 1 while changing the composition of the initially added fraction and the subsequently added fraction of the raw materials as below, to give a resin [A-9] having the following results of the measurement:

<Initially added fraction>

10 parts of acrylonitrile, 10 parts of ethyl acrylate

<Subsequently added fraction>

40 parts of acrylonitrile, 40 parts of ethyl acrylate

<Results of the measurement>

Resin composition: 50% acrylonitrile, 50% ethyl acrylate

Average particle diameter of the resin in the aqueous
5 medium: 0.23 μm

Glass transition temperature: 45°C

Weight average molecular weight: 130,000

[Magnetic material]

10 The kind and the longest length in two-dimensional projection of the magnetic material used in Examples and Comparative Examples of the invention are as follows:

B-1: FL-900 (ferrite) manufactured by Toda Kogyo Corp.,
the longest length of 1.6 μm ,

15 B-2: FH-800 (ferrite) manufactured by Toda Kogyo Corp.,
the longest length of 1.4 μm ,

B-3: Wellmax P-10 (Sm-Co series) manufactured by Sumitomo Metal Industries, Ltd., the maximum length of 5.3 μm ,

20 B-4: Neo40 (Nd-Fe-B series) manufactured by Seiko Sangyo, Co., Ltd., the maximum length of 4.1 μm ,

B-5: SAN40 (Al-Ni-Co series) manufactured by Seiko Sangyo, Co., Ltd., the maximum length of 8.5 μm .

[Examples 1 to 16 and Comparative Examples 1 to 8]

To an aqueous medium wherein any of the resins (A-1 to A-9) obtained in Preparation Examples 1 to 9 was dispersed, any of the magnetic materials (B-1 to B-4) was added in the compositions as shown in [Table 1] to [Table 3], which was stirred for 30 minutes. Then, particle formation by spray drying was carried out using LT-8 which is the spray drying device manufactured by Ohkawara Kakouki Co., Ltd. under the conditions of [Table 1] to [Table 3]. Furthermore, the type of the spray was the disk type, and the collection method was 2 point collection type.

[Table 1]

	Outlet temperature of hot air (°C)	65	65	65	65	65	65	65	65
	Disk rotation No. (rpm)	18,000	18,000	18,000	18,000	18,000	18,000	18,000	18,000
	Total concentration of the resin and the magnetic material in the aqueous medium (% by weight)	46	46	46	46	46	75	56	32
	Productivity in particle formation	AA							
Properties of the composite composition	Average particle diameter (μm)	35	38	33	34	37	36	42	31
	Sphericity	0.85	0.83	0.88	0.84	0.83	0.81	0.86	0.85

[Table 2]

		Example No.							
		9	10	11	12	13	14	15	16
Composition	Kind of the resin	A-1	A-1	A-2	A-3	A-4	A-5	A-6	A-7
	Composition of the resin (part)	100	100	100	100	100	100	100	100
	Kind of the magnetic material	B-1	B-1	B-1	B-1	B-1	B-1	B-1	B-1
	Composition of the magnetic material (part)	100	100	100	100	100	100	100	100
Condition for particle formation	Inlet temperature of hot air (°C)	140	120	120	120	140	140	120	180
	Outlet temperature of hot air (°C)	65	75	65	65	70	70	55	70

	Disk rotation No. (rpm)	18,000	18,000	18,000	18,000	18,000	18,000	18,000	18,000
Total concentration of the resin and the magnetic material in the aqueous medium (% by weight)		46	46	46	46	46	46	46	46
Productivity in particle formation		AA							
Properties of the composite composition	Average particle diameter (μm)	35	36	37	33	35	36	32	33
	Sphericity	0.85	0.83	0.83	0.84	0.81	0.84	0.88	0.92

[Table 3]

the resin and the magnetic material in the aqueous medium (% by weight)								
Productivity in particle formation	AA	AA	BB	CC	CC	BB	CC	CC
Properties of the composite composition	Average particle diameter (μm)	19	21	32	-	-	43	-
	Sphericity	0.42	0.51	0.81	-	-	0.83	-

[Reviews of Examples]

Any of the spherical composite composition of the invention (Examples 1 to 16) is excellent in productivity
 5 and physical properties of the product (the average particle diameter and the sphericity).

On the other hand, when the weight average molecular weight of the resin is more than 1,000,000 (Comparative Examples 1 and 2), the resin is difficult to melt, and thus
 10 formed powders are not bound (bound by the melting of the resin) and the sphericity becomes low. When the glass transition temperature of the resin is lower than 50°C (Comparative Examples 3 and 4), the resin is aggregated or solidified in the disk of the spray parts of the spray dryer,
 15 and thus the productivity is poor. When the inlet temperature of hot air is higher than the temperature which is the glass transition temperature of the resin plus 150°C (Comparative Example 5), the resin is aggregated or solidified in the disk of the spray parts and production is

not possible. When the inlet temperature of hot air in the spray drier is 100°C or less (Comparative Example 6), drying is not enough in the drying chamber of the spray dryer, so the resin contains a lot of water and adheres to the drying 5 chamber, and thus the productivity is poor. In addition, when the amount of the magnetic material to be added is more than 1,000, relative to 100 parts by weight of the resin (Comparative Examples 7 and 8), the produced powders are not bound and the sphericity becomes low.

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Industrial Applicability

The spherical composite composition with high sphericity of the present invention comprising a resin containing unsaturated vinyl units and a magnetic material, 15 can be well used in applications such as a resin magnet, an electric wave absorption material, a magnetic shield material, a magnetic toner material and a toner carrier material. Furthermore, the process of producing a spherical composite composition of the present invention is simple and 20 has good productivity without the need of two or more production steps, whereby it is a process which is very valuable industrially.